

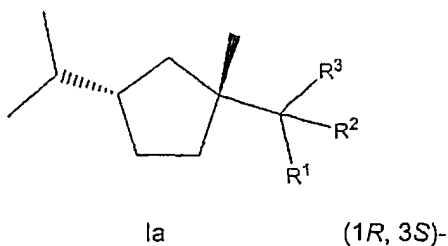
3-ISOPROPYL-1-METHYLCYCLOPENTYL DERIVATIVES AND THEIR USE IN FRAGRANCE APPLICATIONS

The present invention relates to 3-isopropyl-1-methylcyclopentyl derivatives, namely (3-isopropyl-1-methylcyclopentyl)ethanol, (3-isopropyl-1-methylcyclopentyl)ethanone and (3-isopropyl-1-methylcyclopentyl)methanol and their use as fragrances. This invention relates furthermore to a method for their production and to fragrance compositions comprising them.

In the fragrance industry there is a constant demand for new compounds that enhance or improve on odour notes, or impart new odour notes.

It has now been found that certain 3-isopropyl-1-methylcyclopentyl derivatives have much sought-after floral, fruity and woody odour notes, and they are relatively simple and easy to prepare starting from naturally available (1*S*)-(+)- and (1*R*)-(-)-fenchone.

Accordingly, the present invention refers in one of its aspects to the use of a compound of formula Ia and the enantiomer, namely (1*S*,3*R*)- enantiomer, thereof as fragrance



wherein

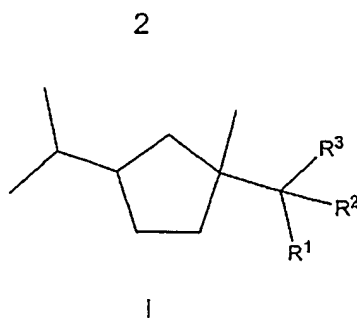
R¹ is hydrogen or methyl;

R² is hydrogen; and

R³ is hydroxyl; or

R² and R³ form together with the carbon atom to which they are attached a carbonyl group.

It has been found that the odour threshold of certain compounds of formula Ia is on an average two times lower than that of the corresponding enantiomer. Accordingly, a compound of formula I



enriched in its (1*R*,3*S*) enantiomer of formula 1a are preferred.

The term "enriched" is used herein to describe a compound having an enantiomeric purity greater than 1:1 in favour of the selected enantiomer. Compounds are preferred having a purity of about 1:3 or greater, e.g. 1:4. Particularly preferred are compounds having an enantiomeric purity of 1:9 or greater, such as 5:95 or 1:99.

Particularly preferred compounds of the present invention are [(1*R*,3*S*)-3-isopropyl-1-methylcyclopentyl]methanol, 1-[(1*R*,3*S*)-3-isopropyl-1-methylcyclopentyl]ethanone, and 1-[(1*R*,3*S*)-3-isopropyl-1-methylcyclopentyl]ethanol.

The compounds according to the present invention may be used alone or in combination with a base material. As used herein, the "base material" includes all known odourant molecules selected from the extensive range of natural products and synthetic molecules currently available, such as essential oils, alcohols, aldehydes and ketones, ethers and acetals, esters and lactones, macrocycles and heterocycles, and/or in admixture with one or more ingredients or excipients conventionally used in conjunction with odourants in fragrance compositions, for example, carrier materials, and other auxiliary agents commonly used in the art.

The following list comprises examples of known odourant molecules, which may be combined with the compounds of the present invention:

- ethereal oils and extracts, e.g. tree moss absolute, basil oil, castoreum, costus root oil, myrtle oil, oak moss absolute, geranium oil, jasmin absolute, patchouli oil, rose oil, sandalwood oil, wormwood oil, lavender oil or ylang-ylang oil;
- alcohols, e.g. citronellol, EbanolTM, eugenol, farnesol, geraniol, Super MuguetTM, linalool, phenylethyl alcohol, SandaloreTM, terpineol or TimberolTM.

- aldehydes and ketones, e.g. α -amylcinnamaldehyde, GeorgywoodTM, hydroxycitronellal, Iso E Super[®], Isoraldeine[®], Hedione[®], maltol, Methyl cedryl ketone, methylionone or vanillin;
- ethers and acetals, e.g. AmbroxTM, geranyl methyl ether, rose oxide or SpirambreneTM.
- esters and lactones, e.g. benzyl acetate, Cedryl acetate, γ -decalactone, Helvetolide[®], γ -undecalactone or Vetivenyl acetate.
- macrocycles, e.g. Ambrettolide, Ethylene brassylate or Exaltolide[®].
- heterocycles, e.g. isobutylcholine.

The compounds of the present invention may be used in a broad range of fragrance applications, e.g. in any field of fine and functional perfumery, such as perfumes, household products, laundry products, body care products and cosmetics. The compounds can be employed in widely varying amounts, depending upon the specific application and on the nature and quantity of other odourant ingredients. The proportion is typically from 0.001 to 20 weight percent of the application. In one embodiment, compounds of the present invention may be employed in a fabric softener in an amount of from 0.001 to 0.05 weight percent. In another embodiment, compounds of the present invention may be used in fine perfumery in amounts of from 0.1 to 20 weight percent, more preferably between 0.1 and 5 weight percent. However, these values are given only by way of example, since the experienced perfumer may also achieve effects or may create novel accords with lower or higher concentrations.

The compounds of the present invention may be employed into the fragrance application simply by directly mixing the fragrance composition with the fragrance application, or they may, in an earlier step be entrapped with an entrapment material, for example, polymers, capsules, microcapsules and nanocapsules, liposomes, film formers, absorbents such as carbon or zeolites, cyclic oligosaccharides and mixtures thereof, or they may be chemically bonded to substrates, which are adapted to release the fragrance molecule upon application of an external stimulus such as light, enzyme, or the like, and then mixed with the application.

Thus, the invention additionally provides a method of manufacturing a fragrance application, comprising the incorporation of a compound of formula I enriched in one of their enantiomers, as a fragrance ingredient, either by directly admixing the compound to the application or by admixing a fragrance composition comprising a compound of formula I enriched in one of their enantiomers, which may then be mixed to a fragrance application, using conventional techniques and methods.

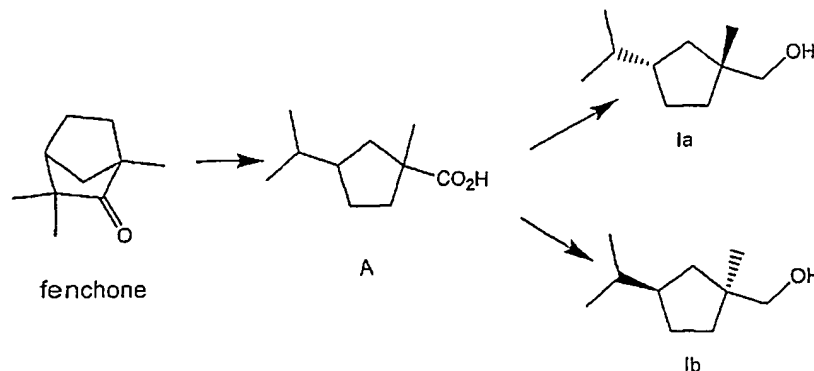
As used herein, "fragrance application" means any product, such as fine perfumery, e.g. perfume and eau de toilette; household products, e.g. detergents for dishwasher, surface cleaner; laundry products, e.g. softener, bleach, detergent; body care products, e.g. shampoo, shower gel; and cosmetics, e.g. deodorant, vanishing creme, comprising an odourant. This list of products is given by way of illustration and is not to be regarded as being in any way limiting.

Compared to most odourant molecules known in the art having floral odor properties, such as hydroxycitronellal, geranol, linalool and 4-(4-hydroxy-4-methylpentyl)cyclohex-3-ene-1-carbaldehyde, the compounds of formula Ia of the present invention, wherein R^3 is hydroxyl, are exceptionally stable both, under basic and acidic conditions, thus making them particularly useful for a large variety of fragrance applications.

Compounds of formula Ia and the enantiomers thereof may be prepared by the Haller-Bauer rearrangement of (1*R*)-(-)-fenchone / (1*S*)-(+)-fenchone (1,3,3-trimethyl-2-norbornanone) followed by hydrolysis to 3-isopropyl-1-methylcyclopentanecarboxylic acid under alkali conditions, e.g. in the presence of a base such as NaOH or KOH. The resulting acid will then be reacted with methyllithium to give a compound of formula I wherein R^2 and R^3 form together with the carbon atom to which they are attached a carbonyl group. To give further compounds of the present invention, the resulting ketone may be transformed to a secondary alcohol through reduction, e.g. with NaBH₄.

(3-Isopropyl-1-methylcyclopentyl)methanol may be prepared by reduction of 3-isopropyl-1-methylcyclopentanecarboxylic acid (A), which has been prepared by rearrangement of fenchone, in the presence of LAH to the corresponding alcohol, as shown in scheme 1.

Scheme 1:



Optically pure compounds of formula 1a and 1b and enantiomeric mixtures of a compound of formula 1 enriched in one of the enantiomers, i.e. a compound of formula 1a or 1b, may be synthesised, starting from optically pure fenchone or an enantiomeric mixture enriched in either (1*R*)-(-)-fenchone or (1*S*)-(+)-fenchone.

The invention is now further described with reference to the following non-limiting examples.

All end products described in the following Examples 1 to 6 are colourless oils. They were obtained starting from (1*R*)-(-)- and (1*S*)-(+)-fenchone that contained 8% and 2% respectively of the other enantiomer. The reported NMR data were measured under the following general conditions: ¹H at 400 and ¹³C at 100 MHz; in CDCl₃; chemical shifts (δ) in ppm downfield from TMS; coupling constants *J* in Hz.

Example 1: [(1*R*,3*S*)-3-Isopropyl-1-methylcyclopentyl]methanol

A solution of (1*R*,3*S*)-3-Isopropyl-1-methylcyclopentanecarboxylic acid (70.0 g, 0.41 mol), obtained from (1*R*)-(-)-fenchone (V. Braun, J.; Jacob, A. *Chem. Ber.* 1933, 66, 1461) in diethyl ether (100 ml) was slowly added, under nitrogen, to a suspension of lithium aluminium hydride (13.3 g, 0.35 mol) in the same solvent (500 ml). After heating at reflux during 3 h, the reaction mixture was cooled down to 10°C, 2N NaOH solution (70 ml) was carefully added and stirring continued for 0.5 h. The white solid was filtered off, the filtrate washed with brine (2 x 500 ml), dried (MgSO₄) and concentrated *in vacuo*. The crude product (79.0 g) was purified by distillation using a 10 cm Vigreux column (0.9-1.1 mbar, 96-98°C) to give [(1*R*,3*S*)-3-isopropyl-1-methylcyclopentyl]methanol (57.0 g, 90% yield).

$^1\text{H-NMR}$: δ 0.87 (*d*, $J = 6.7$, 3H), 0.88 (*d*, $J = 6.7$, 3H), 1.01 (*s*, 3H), 1.08 (*dd*, $J = 12.3$, 11.0, 1H), 1.16-1.38 (*m*, 3H), 1.48 (*ddd*, $J = 12.4$, 6.9, 0.8, 1H), 1.53-1.72 (*m*, 3H), 1.74-1.87 (*m*, 1H), 3.35 (*d*, $J_{AB} = 10.4$, 1H), 3.39 (*d*, $J_{AB} = 10.4$, 1H). $^{13}\text{C-NMR}$: δ 21.5 (2q), 25.0 (q), 30.4 (t), 33.8 (d), 35.6 (t), 41.5 (t), 43.8 (s), 46.9 (d), 72.1 (t). $[\alpha]_{\text{D}}^{22} -12.0$ (c 1.0, EtOH).

Odour description: floral, green, jasmine, lily-of-the-valley, fresh, clean.

Example 2: [(1*S*,3*R*)-3-Isopropyl-1-methylcyclopentyl]methanol

Prepared according to the experimental procedure of Example 1 starting from (1*S*)-(+)-fenchone.

$[\alpha]_{\text{D}}^{22} +13.5$ (c 1.0, EtOH).

Odour description: floral, fruity, green, rosy, hesperidic (grapefruit).

Example 3: 1-[(1*R*,3*S*)-3-Isopropyl-1-methylcyclopentyl]ethanone

A 1.6M solution of methyllithium in diethyl ether (200 ml, 0.32 mol) was added dropwise during 25 min. into a solution of (1*R*,3*S*)-3-Isopropyl-1-methylcyclopentanecarboxylic acid (25.5 g, 0.15 mol) in THF (250 ml) at 0°C. After stirring at 0°C for 3 h, chlorotrimethylsilane (151 ml, 1.2 mol) was added with cooling and the reaction mixture was allowed to warm up to room temperature, poured on ice-cold water (200 ml), stirred for 0.5 h and extracted with MTBE (2 x 250 ml). The combined organic phases were washed with water (200 ml), 2M NaOH (150 ml) and brine (3 x 200 ml), dried (MgSO_4) and concentrated in vacuo to give the crude 1-[(1*R*,3*S*)-3-isopropyl-1-methylcyclopentyl]ethanone (27.6 g), a sample of which (1.5 g) was purified by bulb-to-bulb distillation (0.93 g, 68% yield).

$^1\text{H-NMR}$: δ 0.89 (2*d*, $J = 6.6$, 6H), 1.19 (*s*, 3H), 1.24 (*dq*, $J = 12.4$, 9.1, 1H), 1.34-1.43 (*m*, 2H), 1.56-1.77 (*m*, 3H), 1.81-1.90 (*m*, 1H), 2.09 (*ddd*, $J = 13.1$, 9.1, 4.0, 1H), 2.15 (*s*, 3H). $^{13}\text{C-NMR}$: δ 21.3 (q), 21.4 (q), 25.0 (q), 25.3 (q), 30.2 (t), 33.3 (d), 35.6 (t), 41.0 (t), 46.6 (d), 55.3 (s), 213.0 (s). $[\alpha]_{\text{D}}^{22} -1.0$ (c 1.1, EtOH).

Odour description: earthy/mossy, green, woody.

Example 4: 1-[(1S,3R)-3-Isopropyl-1-methylcyclopentyl]ethanone

Prepared according to the experimental procedure of Example 3 starting from (1S)-(+)-fenchone.

$[\alpha]_D^{22} +1.0$ (c 1.1, EtOH).

Odour description: floral, agrestic, fruity, green.

Example 5: 1-[(1R,3S)-3-Isopropyl-1-methylcyclopentyl]ethanol

A solution of 1-[(1R,3S)-3-isopropyl-1-methylcyclopentyl]ethanone from Example 3 (3.0 g, 18 mmol) in ethanol (8 ml) was added to a cold (ice-bath) solution of sodium borohydride (0.42 g, 10.7 mmol) in the same solvent (17 ml). After 1.5 h stirring at room temperature, the reaction mixture was poured on ice-cold 2M HCl (100 ml) and extracted with MTBE (2 x 100 ml). The combined organic phases were washed with brine (2 x 50 ml), dried (MgSO₄) and concentrated in vacuo. The crude product (2.8 g) was purified by bulb-to-bulb distillation (2.34 g, 77% yield, diastereoisomer ratio ~1:1).

¹H-NMR: δ 0.87 (d, J = 6.6, 3H), 0.875 (d, J = 6.6, 3H), 0.88 (2d, J = 6.6, 6H), 0.92 (s, 3H), 0.93 (s, 3H), 1.05 (t, J = 11.7, 1H), 1.12 (d, J = 6.4, 3H), 1.125 (d, J = 6.4, 3H), 1.14 (t, J = 11.8, 1H), 1.17-1.74 (m, 12H), 1.47 (2s, 2H), 1.78-1.88 (m, 2H), 3.53 (q, J = 6.3, 1H), 3.55 (q, J = 6.3, 1H). ¹³C-NMR: δ 18.5 (2q), 21.3 (2q), 21.4 (3q), 21.5 (q), 29.8 (t), 30.0 (t), 33.7 (2d), 35.8 (t), 35.9 (t), 41.9 (2t), 46.3 (2d), 46.8 (s), 46.9 (s), 75.4 (d), 75.7 (d). $[\alpha]_D^{22} -7.0$ (c 1.0, EtOH).

Odour description: floral, earthy/mossy, slightly terpineol/earthy.

Example 6: 1-[(1S,3R)-3-Isopropyl-1-methylcyclopentyl]ethanol

Prepared according to the experimental procedure of Example 5 starting from (1S)-(+)-fenchone.

$[\alpha]_D^{22} +8.0$ (c 1.0, EtOH).

Odour description: hesperidic/citrus, fruity, green, fresh (grapefruit, rhubarb).

Example 7: Feminine Fine Fragrance

<u>Ingredient</u>	<u>Parts per weight</u>
Citronellol	50
Cyclamen aldehyde	15
Diethyl malonate	5
Dipropylene glycol (DPG)	149
Florhydral	12
Gardenol	10
Geraniol	50
Hedione	25
alpha-Hexylcinnamaldehyde	200
Hydroxycitronellal	35
Isocyclocitral 1% in DPG	15
Isojasmone	2
Jasmal	40
Jasmonyl	20
Lemon oil	10
Lilial	25
Linalool	65
Linalyl acetate	50
Methyl diantilis	2
Petitgrain Paraguay oil	5
Phenethyl alcohol	65
Silvial	100
<u>[(1R,3S)-3-Isopropyl-1-methylcyclopentyl]methanol</u>	<u>50</u>
Total	1000

* for chemical names see Flavor and Fragrance Materials – 2003, Allured Publishing Corp. Carol Stream Ill., U.S.A..

The presence of 5% of [(1*R*,3*S*)-3-Isopropyl-1-methylcyclopentyl]methanol confers to this formula a creamy, lily-of-the-valley aspect.

Example 8: Floral Composition for Soap

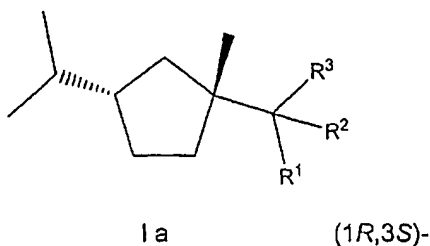
<u>Ingredient</u>	<u>Parts per weight</u>
Agrumex	100
Benzophenone	60
Benzyl acetate	55
Bergamot base	80
4- <i>t</i> -Butylcyclohexyl acetate	150
Diphenyl oxide	20
Dipropylene glycol (DPG)	78
Ebanol	20
Hydroxycitronellal	200
Jasmine base	80
Methyl Phenylacetate	2
Nerol	20
Phenylpropyl alcohol	40
Rose base	100
Rhodinol	65
Sandela	30
Silvial	100
<u>[(1<i>R</i>,3<i>S</i>)-3-Isopropyl-1-methylcyclopentyl]methanol</u>	<u>50</u>
Total	1250

* for chemical names see Flavor and Fragrance Materials – 2003, Allured Publishing Corp. Carol Stream Ill., U.S.A..

[(1*R*,3*S*)-3-Isopropyl-1-methylcyclopentyl]methanol makes this lily-of-the-valley fragrance velvety and rich.

Claims

1. The use of a compound of formula Ia and the enantiomer thereof as fragrance,



wherein

R^1 is hydrogen or methyl;

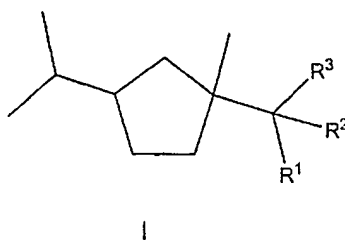
R^2 is hydrogen; and

R^3 is hydroxyl; or

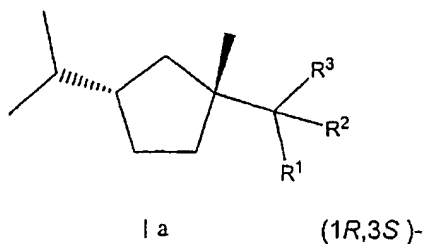
R^2 and R^3 form together with the carbon atom to which they are attached a carbonyl group.

2. The use as fragrance of a compound according to claim 1 selected from the group consisting of [(1*R*,3*S*)-3-isopropyl-1-methylcyclopentyl]methanol, [(1*S*,3*R*)-3-isopropyl-1-methylcyclopentyl]methanol, 1-[(1*R*,3*S*)-3-isopropyl-1-methylcyclopentyl]ethanone, 1-[(1*S*,3*R*)-3-isopropyl-1-methylcyclopentyl]ethanone, 1-[(1*R*,3*S*)-3-isopropyl-1-methylcyclopentyl]ethanol and 1-[(1*S*,3*R*)-3-isopropyl-1-methylcyclopentyl]ethanol.

3. The use as fragrance of a compound of formula I



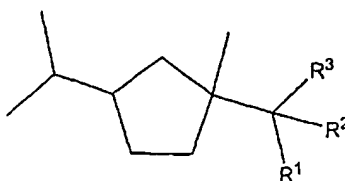
enriched in the enantiomer having the formula Ia



wherein R^1 , R^2 and R^3 have the same meaning as given in claim 1.

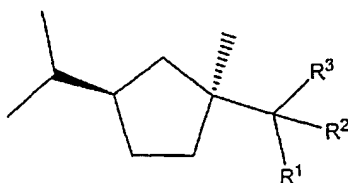
11

The use as fragrance of a compound of formula I



I

enriched in the enantiomer having the formula Ib



Ib

(1*S*,3*R*)-

wherein R^1 , R^2 and R^3 have the same meaning as given in claim 1.

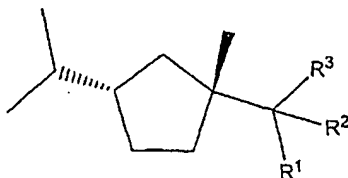
The use of a compound as defined in one of the preceding claims in fragrance applications.

A fragrance application comprising a compound as defined in any of the preceding claims 1 - 4.

A fragrance application according to claim 6 wherein the fragrance application is a perfume, household product, laundry product, body care product or cosmetic products.

A method of manufacturing a fragrance application, comprising the step of incorporating a compound of formula Ia or its enantiomer as defined in claim 1, 2, 3 and 4.

A compound of formula Ia



Ia

(1*R*,3*S*)-

wherein

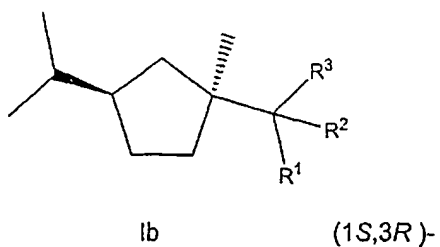
R^1 is hydrogen or methyl;

R^2 is hydrogen; and

R^3 is hydroxyl; or

R^2 and R^3 form together with the carbon atom to which they are attached a carbonyl group.

. A compound of formula Ib



wherein

R^1 is hydrogen or methyl;

R^2 is hydrogen; and

R^3 is hydroxyl; or

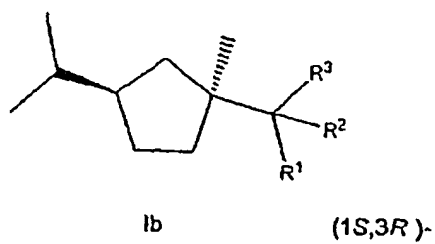
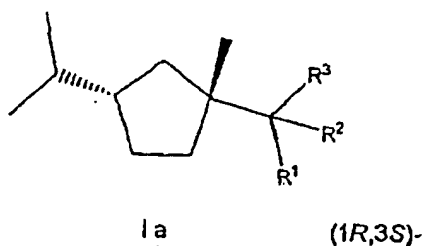
R^2 and R^3 form together with the carbon atom to which they are attached a carbonyl group.

INTERNATIONAL SEARCH REPORT

PCT/CH2004/000604

Box No. IV Text of the abstract (Continuation of item 5 of the first sheet)

This invention relates to 3-isopropyl-1-methylcyclopentyl derivatives of formula Ia and Ib, where R^1 , R^2 and R^3 are as defined in the claims, and their use in fragrance applications.



A. CLASSIFICATION OF SUBJECT MATTER

IPC 7 C11B9/00 C07C31/135 C07C49/297

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC 7 C11B C07C

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

EPO-Internal, BEILSTEIN Data, CHEM ABS Data, WPI Data, PAJ

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	PATENT ABSTRACTS OF JAPAN vol. 0151, no. 45 (C-0823), 12 April 1991 (1991-04-12) & JP 3 024198 A (TAKASAGO INTERNATIONAL), 1 February 1991 (1991-02-01) abstract	1-10
A	US 4 234 463 A (M.A. SPRECKER, ET AL.) 18 November 1980 (1980-11-18) claim 1	1-10
A	US 4 533 492 A (R.J. TOKARZEWSKI, ET AL.) 6 August 1985 (1985-08-06) claim 1	1-10

☐ Further documents are listed in the continuation of box C.☒ Patent family members are listed in annex.

* Special categories of cited documents :

A document defining the general state of the art which is not considered to be of particular relevance

E earlier document but published on or after the international filing date

L document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)

O document referring to an oral disclosure, use, exhibition or other means

P document published prior to the international filing date but later than the priority date claimed

T later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention

X document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone

Y document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.

G document member of the same patent family

Date of the actual completion of the international search

10 December 2004

Date of mailing of the international search report

07/01/2005

Name and mailing address of the ISA

European Patent Office, P.B. 5818 Patentlaan 2
NL - 2280 HV Rijswijk
Tel. (+31-70) 340-2040, Tx. 31 651 epo nl,
Fax: (+31-70) 340-3016

Authorized officer

English, R

Patent document cited in search report		Publication date	Patent family member(s)	Publication date
JP 3024198	A	01-02-1991	JP 2065384 C	24-06-1996
			JP 7091554 B	04-10-1995
US 4234463	A	18-11-1980	US 4195100 A	25-03-1980
			US 4206092 A	03-06-1980
			US 4195099 A	25-03-1980
			US 4222941 A	16-09-1980
			US 4256121 A	17-03-1981
			US 4197328 A	08-04-1980
			US 4263149 A	21-04-1981
			US 4217371 A	12-08-1980
			US 4207214 A	10-06-1980
			US 4231940 A	04-11-1980
			US 4208308 A	17-06-1980
			US 4269862 A	26-05-1981
			US 4247423 A	27-01-1981
			US 4250342 A	10-02-1981
US 4533492	A	06-08-1985	US 4449009 A	15-05-1984